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Short presentation

Older adults are the main consumers of drugs but are rarely included in pivotal randomized clinical trials. Hence, post-marketing surveillance is a pivotal activity to better characterize the safety and effectiveness of medications in older patients.

I am currently coordinating a team integrating multidisciplinary expertise from Pharmacoepidemiology, Pharmacovigilance, and Data science to catalyze the identification of subgroups of older patients at higher risk of adverse drug reactions or not sufficiently benefitting from pharmacological treatments using post-marketing data sources (e.g., administrative and spontaneous reporting databases). The team has a strong translational perspective as its final goal is to promote the integration of discoveries into patient care.

The team has access to the biggest cohort of older individuals in Denmark covering the entire Danish population aged ≥ 65 for which they have socio-demographic information, hospital- and ambulatory access to Danish hospitals, pharmacological and non-pharmacological treatments, laboratory results (i.e., biochemical, genetic, and immunological), surgery, and procedures.

Qualifications

Regulatory affairs, Certified program in EU Regulatory Affairs
Award Date: 1 Sep 2017

Translational pharmacology, PhD
Award Date: 20 Dec 2016

Employment

Associate Professor

Pharmaceutical Informatics

København Ø

1 May 2018 → nu

Assistant professor, tenure track

Pharmacotherapy

København Ø, Denmark

1 Sep 2019 → 3 Jan 2020

Research outputs

Artificial intelligence for the optimal management of community-acquired pneumonia

Barbieri, M. A., Battini, V. & Sessa, Maurizio, 2024, In: Current Opinion in Pulmonary Medicine. 30, 3, p. 252-257

Comparing major and mild cognitive impairment risks in older type-2 diabetic patients: a Danish register-based study on dipeptidyl peptidase-4 inhibitors vs. glucagon-like peptide-1 analogues

Battini, V., Barbieri, M. A., Carnovale, C., Spina, E., Clementi, E. & Sessa, Maurizio, 2024, In: Journal of Neurology. 271, p. 3417-3425

Drug Repurposing in Crohn's Disease Using Danish Real-World Data

Shakibfar, Saeed, Allin, K. H., Jess, T., Barbieri, M. A., Battini, V., Simoncic, E., Kirchgessner, J., Ulven, Trond & Sessa, Maurizio, 2024, In: Pragmatic and observational research. 15, p. 17-29

Exploring the impact of co-exposure timing on drug-drug interactions in signal detection through spontaneous reporting system databases: a scoping review

Cocco, M., Carnovale, C., Clementi, E., Barbieri, M. A., Battini, V. & Sessa, Maurizio, 2024, In: Expert Review of Clinical Pharmacology. 17, 5-6, p. 441-453

Impact of the early COVID-19 pandemic on adult mental health-related dispensed medications, hospitalizations and specialist outpatient visits in Norway and Sweden: Interrupted time series analysis

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New data-driven method to predict the therapeutic indication of redeemed prescriptions in secondary data sources: a case study on antiseizure medications users aged ≥ 65 identified in Danish registries

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Persistence and adherence with Latanoprost: A Danish register-based cohort study in older patients with glaucoma

Amiri, D., Sessa, Maurizio, Andersen, Morten & Kolko, Miriam, 2024, In: Acta Ophthalmologica. 102, 2, p. 172-178 7 p.

Ticagrelor or prasugrel vs. clopidogrel in patients with atrial fibrillation undergoing percutaneous coronary intervention for myocardial infarction

Godtfredsen, S. J., Kragholm, K. H., Kristensen, A. M. D., Bekfani, T., Sørensen, Rikke, Sessa, Maurizio, Torp-Pedersen, Christian, Bhatt, D. L. & Pareek, M., 2024, In: European Heart Journal Open. 4, 1, 9 p., oead134.

Timing Matters: A Machine Learning Method for the Prioritization of Drug-Drug Interactions Through Signal Detection in the FDA Adverse Event Reporting System and Their Relationship with Time of Co-exposure

Battini, V., Cocco, M., Barbieri, M. A., Powell, G., Carnovale, C., Clementi, E., Bate, A. & Sessa, Maurizio, 2024, (E-pub ahead of print) In: Drug Safety.

A novel approach for pharmacological substantiation of safety signals using plasma concentrations of medication and administrative/healthcare databases: a case study using Danish registries for an FDA warning on lamotrigine

Wang, W., Battini, V., Carnovale, C., Noordam, R., van Dijk, K. W., Kragholm, K. H., van Heemst, D., Soeorg, H. & Sessa, Maurizio, 2023, In: Pharmacological Research. 193, 8 p., 106811.

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Al-Azzawi, F., Mahmoud, I., Haguinet, F., Bate, A. & Sessa, Maurizio, 2023, In: Drug Safety. 46, p. 743-751

Machine learning-driven development of a disease risk score for COVID-19 hospitalization and mortality: a Swedish and Norwegian register-based study

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Ubrogepant and rimegepant: systematic review, meta-analysis, and meta-regression of clinical studies

Dong, G., Kjærgaard, N. A., Shakibfar, Saeed & Sessa, Maurizio, 2023, In: *Expert Opinion on Drug Safety*. 22, 1, p. 59-70

Ubrogepant and rimegepant: signal detection using spontaneous reports of adverse events from the Food and Drug Administration Adverse Event Reporting System

Battini, V., Carnovale, C., Clementi, E. & Sessa, Maurizio, 2023, In: *Expert Opinion on Drug Safety*. 22, 11, p. 1105-1112

Venous thromboembolism with use of hormonal contraception and non-steroidal anti-inflammatory drugs: nationwide cohort study

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A machine-learning guided method for predicting add-on and switch in secondary data sources: A case study on anti-seizure medications in Danish registries

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Pharmacological and epidemiological considerations while constructing treatment episodes using observational data: A simulation study

Pazzagli, L., Andersen, Morten & Sessa, Maurizio, 2022, In: *Pharmacoepidemiology and Drug Safety*. 31, 1, p. 55-60

Pharmacom-epi: A framework for integrating pharmacometric modeling into Pharmacoepidemiological research using real-world data

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Rationale and performances of a data-driven method for computing the duration of pharmacological prescriptions using secondary data sources

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The PHARMACOM-EPI framework for integrating pharmacometric modelling into pharmacoepidemiological research using real-world data: application to assess death associated with valproate

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The Relationship Between Valproate and Lamotrigine/Levetiracetam Use and Prognosis in Patients With Epilepsy and Heart Failure: A Danish Register-Based Study

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A systematic review, meta-analysis and meta-regression evaluating the adverse reactions to erenumab in the preventive treatment of migraine

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Detecting deviations from the efficacy and safety results of single-arm trials using real-world data: The case of a CAR-T cell therapy in B-cell lymphoma

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New Insight on the Safety of Erenumab: An Analysis of Spontaneous Reports of Adverse Events Recorded in the US Food and Drug Administration Adverse Event Reporting System Database

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Pharmacoepidemiological methods for computing the duration of pharmacological prescriptions using secondary data sources

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The relationship between valproate versus lamotrigine/levetiracetam use and prognosis in patients with epilepsy and heart failure: A Danish register-based study

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